Claims.

1. An 1-[(indol-3-yl)carbonyl]piperazine derivative having the general formula I

Formula I

wherein

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R represents 1-4 substituents independently selected from H,  $(C_{1-4})$ alkyl (optionally substituted with halogen),  $(C_{1-4})$ alkyloxy (optionally substituted with halogen), halogen, OH, NH<sub>2</sub>, CN and NO<sub>2</sub>;

10  $R_1$  is  $(C_{5-8})$  cycloalkyl or  $(C_{5-8})$  cycloalkenyl;

R<sub>2</sub> is H, methyl or ethyl;

 $R_3$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_5$  and  $R_6$  are independently hydrogen or ( $C_{1-4}$ )alkyl, optionally substituted with ( $C_{1-4}$ )alkyloxy, halogen or OH;

 $R_6$  is hydrogen or  $(C_{1-4})$ alkyl, optionally substituted with  $(C_{1-4})$ alkyloxy, halogen or

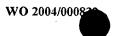
15 OH; or

 $R_6$  forms together with  $R_7$  a 4-7 membered saturated heterocyclic ring, optionally containing a further heteroatom selected from O and S;

 $R_7$  forms together with  $R_6$  a 4-7 membered saturated heterocyclic ring, optionally containing a further heteroatom selected from O and S; or

- 20  $R_7$  is H,  $(C_{1-4})$ alkyl or  $(C_{3-5})$ cycloalkyl, the alkyl groups being optionally substituted with OH, halogen or  $(C_{1-4})$ alkyloxy; or
  - a pharmaceutically acceptable salt thereof.
- 2. The 1-[(indol-3-yl)carbonyl]piperazine derivative of claim 1, wherein R<sub>2</sub> is H and R<sub>1</sub> is (C<sub>5-6</sub>)cycloalkyl.
  - 3. The 1-[(indol-3-yl)carbonyl]piperazine derivative of claim 2, wherein R is (C<sub>1-4</sub>)alkyloxy or halogen.

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- 4 The 1-[(indol-3-yl)carbonyl]piperazine derivative of claim 3, wherein R represents a methoxy group at the 7-position of the indole ring.
- 5. The 1-[(indol-3-yl)carbonyl]piperazine derivative of claim 4, wherein R<sub>3</sub>, R<sub>3</sub>', R<sub>4</sub>', R<sub>5</sub>, R<sub>5</sub>' and R<sub>6</sub>' are H; R<sub>4</sub>, R<sub>6</sub> and R<sub>7</sub> are independently H or (C<sub>1-4</sub>)alkyl; or R<sub>6</sub> forms together with R<sub>7</sub> a 5- or 6-membered saturated heterocyclic ring and R<sub>4</sub> is H or (C<sub>1-4</sub>)alkyl.
- 6. The 1-[(indol-3-yl)carbonyl]piperazine derivative according to formula I of claim 1 which is selected from:
  - 1-{[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl}-3,5-dimethyl 4-ethylpiperazine;
  - 1-{[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl}-3,4,5-tri-methylpiperazine;
- (S)-1-{[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl}-3,4-dimethylpiperazine;
  - (S)-2-{[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl}-octahydro-2*H*-pyrido-[1, 2-a]pyrazine;
  - (S)-2-{[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl}-octahydro-2*H*-pyrrolo-[1, 2-a]pyrazine; and
  - (S)-2-{[1-(cyclopentylmethyl)l-7-methoxy-1*H*-indol-3-yl]carbonyl}-octa-hydro-2*H*-pyrido-[1, 2-a]pyrazine;

or a pharmaceutically acceptable salt thereof.

- 7. The 1-[(indol-3-yl)carbonyl]piperazine derivative of any one of claims 1-6 for use in therapy.
- 8. A pharmaceutical composition comprising an 1-[(indol-3-yl)carbonyl]piperazine derivative of any one of claims 1-6 together with a pharmaceutically acceptable carrier therefor.
  - 9. Use of an 1-[(indol-3-yl)carbonyl]piperazine derivative of formula I as defined in claim 1, in the preparation of a medicament for the treatment of pain.

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